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<p><b>(54) Title:</b> PROTEINS INVOLVED IN THE REGULATION OF CELL-CELL ADHESION AND USES THEREOF</p> <p><b>(57) Abstract</b></p> <p>A p100 protein, which has a molecular weight of about 100 kDa and appears to be related to p120, is involved in the regulation of cell-cell adhesion.</p>			

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**PROTEINS INVOLVED IN THE REGULATION OF CELL-CELL ADHESION AND USES THEREOF**

The present invention relates to certain proteins which may be involved in regulating physiological changes (e.g. changes in cell-cell adhesion) and to uses thereof.

Cell adhesion is important for a wide variety of regulatory and developmental processes. The cadherins comprise a family of transmembrane, cell surface glycoproteins that mediate  $\text{Ca}^{2+}$ -dependent cell-cell adhesion in a homotypic manner (Takeichi, 1991). In cells with well developed intercellular junctions, the cadherins are localized to the adherens junction (Boller et al., 1985) but appear to influence other intercellular junctions such as gap junctions (Matsuzaki et al., 1990; Musil et al., 1990) and tight junctions (Gumbiner and Simons, 1986; Gumbiner et al., 1988). The adherens junction also plays a crucial role in the development and maintenance of cell polarity (see Nelson, 1992) and its dysfunction has been strongly implicated in the invasiveness and carcinogenesis of tumour cells (see e.g. Behrens et al., 1989; Frixen et al., 1991; Vleminckx et al., 1991; Shimoyama et al., 1992; and Hedrick et al., 1993; Tsukita et al., 1993; Birchmeier and Behrens, 1994).

The conserved cytoplasmic domain of cadherins is known to associate with three proteins, termed  $\alpha$ -,  $\beta$ - and  $\gamma$ -catenin (Ozawa et al., 1989), which serve to link cadherins to the actin-based cortical cytoskeleton (Hirano et al., 1987). The association of cadherins with catenins is essential for intercellular  $\text{Ca}^{2+}$ -dependent adhesiveness (Nagafuchi and Takeichi, 1988; Ozawa et al., 1990; Kintner, 1992).  $\alpha$ -catenin is homologous to vinculin (Herrenknecht et al., 1991; Nagafuchi et al., 1991), making it a good candidate for interaction with the actin-based cytoskeleton (see Ozawa et al., 1990; Hirano et al., 1992).  $\beta$ -catenin is homologous to the

Drosophila segment polarity gene *armadillo*, suggesting a role in developmental signalling in vertebrates (McCrea et al., 1991).  $\gamma$ -catenin is probably identical to plakoglobin (Knudsen and Wheelock, 1992; but see Piepenhagen and Nelson, 1993), which again is homologous to *armadillo* (see Franke et al., 1989; Peifer and Wieschaus, 1990). Indeed,  $\beta$ -catenin and plakoglobin appear to form a multigene family (Peifer et al., 1992).

A repeating 42 amino acid motif that was originally identified in *armadillo* (Riggieman et al., 1989) has also been found in several other proteins, including  $\beta$ -catenin and plakoglobin, with a variety of functions (Peifer et al., 1994). These include the APC gene product, a tumour suppressor protein (Kinzler et al., 1991), p120, a  $\text{pp}_60^{\text{src}}$  substrate (Reynolds et al., 1992), smgGDS, an exchange factor for ras-related G proteins (Kikuchi et al., 1992), a suppressor of RNA polymerase I mutations in yeast (Yano et al., 1992; 1994) and band 6 protein, a major desmosomal constituent (Hatzfield et al., 1994). The function of the repeats in these arm proteins is unknown. Interestingly, the APC gene product associates with  $\beta$ -catenin (Rubinfeld et al., 1993; Su et al., 1993), supporting an important role for catenins in intracellular processes that regulate cell growth. Furthermore, this illustrates that cacherins are not exclusive cellular partners of catenins, raising the possibility of other interactions among catenins, cacherins and arm proteins, important in a variety of biological processes.

p120 was initially identified as one of several substrates of the tyrosine kinase  $\text{pp}_60^{\text{src}}$  (Reynolds et al., 1989; Kanner et al., 1990). It is membrane-associated and can be myristoylated, but does not appear to be glycosylated (Kanner et al., 1991). Mutational analysis suggested that tyrosine phosphorylation of p120 is necessary for  $\text{pp}_60^{\text{src}}$ -mediated cellular transformation (Linder and

Burr, 1988; Reynolds et al., 1989). Tyrosine phosphorylation of p120 was also observed in response to epidermal growth factor, platelet-derived growth factor, colony-stimulating factor 1 and in polyoma virus middle T antigen-transformed cells (Dowing and Reynolds, 1991; Kanner et al., 1991), but the exact role of p120 in cellular physiology and pathology was not clear.

According to the present invention, there is provided a protein having a molecular weight of about 100 kDa which is associated in vivo with the cadherin/catenin complex of epithelial or endothelial cells (e.g. by binding). This protein is referred to herein as "p100". p100 is immunologically related to p120 by virtue of cross-reactivity of antibody to p120 with p100.

Partial amino acid sequence data of human p100 is given in Fig. 9 where it is compared with sequence data already available for mouse p120. The present invention covers the specific protein of molecular weight of about 100 kDa described in the present examples and other proteins having substantial amino-acid homology therewith (excluding p120, and optionally also excluding proteins having greater amino acid homology with p120 than with p100). Human p100 is particularly preferred.

The term "substantial amino acid homology" is used herein to cover proteins having at least 50%, preferably more than 90% or more than 95% amino acid homology with another protein. The present invention also covers fragments of these proteins or of p100 itself. Preferably these fragments are at least twenty amino acids long, more preferably they are at least fifty or at least one hundred amino acids long. These proteins or fragments may be partially or totally tyrosine phosphorylated. They may be provided in glycosylated or non-glycosylated form. Desirably they are provided in

substantially pure form. One definition of "substantially pure form" is a form which is substantially free of other proteins.

5 The present invention also includes nucleic acid sequences (preferably DNA sequences) coding for the above mentioned proteins or fragments. These sequences may be in isolated form. They may be incorporated as part of a recombinant nucleic acid molecule e.g. as part of a vector. The vector may be incorporated into a host cell  
10 and used for expression of p100.

Nucleic acid sequences complementary to the aforesaid sequences may be useful in antisense studies to alter the expression of gene products. Such sequences  
15 are therefore also within the scope of the present invention.

In view of the data provided herein, it is believed that the protein p100 may be involved in the regulation  
20 of cell tight junction permeability. Phosphorylation of one or more tyrosine residues of p100 may be involved in increasing tight junction permeability and dephosphorylation of one or more tyrosine residues may be involved in decreasing tight junction permeability. The above comments also apply in respect of p120, which the  
25 present inventors have shown to be immunologically closely related to p100.

p100 and p120 are therefore useful for studying cell-cell adhesion generally and tight junction permeability in particular. They may therefore be used, for example, to investigate the effect of tyrosine kinase and/or of tyrosine phosphatase on tight junction permeability. Variants of these proteins may be prepared in order to investigate which regions of p100 or p120 are important in regulating tight junction permeability.  
30

Our data indicate that p120 and p100 associate with

the cadherin/catenin complex. Furthermore, since p120 and p100 are substrates for tyrosine kinases, it follows that p120 and p100 proteins per se and the phosphorylation of these proteins may influence cellular functions directly, and indirectly, regulated by cadherins and catenins. Such an influence could be mediated by the physical association between p120, p100 and the cadherin/catenin complex, in which case modulation could be achieved by enhancing or blocking the expression of p120 and p100. Disruption of the association could also be achieved, for example, by small molecule mimetics of the site of binding. As indicated above, the function of p120 and p100 may also be regulated by their tyrosine phosphorylation, in which case appropriate modulation of kinases and phosphatases could modulate the function of cadherins and other cadherin-dependent processes.

p100 and p120 may be of utility in studying tumour cell invasion and metastasis, breast cancer, colorectal cancer, gastric carcinoma, gynaecological cancer, lung cancer, oesophageal cancer, prostate cancer, scirrhous cancer, tissue regeneration, for example. Such studies could lead to diagnostic methods or treatments of these disorders.

The *Drosophila* gene product *armadillo* is homologous to  $\beta$ -catenin and has been strongly implicated in developmental processes. It is therefore possible that p120 and p100, as  $\beta$ -catenin-related proteins that interact with cadherins, may also be involved in developmental defects and abnormalities.

The present invention also provides an agent which alters or blocks association between p100 or p120 with the catenin/cadherin complex of epithelial or endothelial cells for use in the treatment or diagnosis of a disease.

5           The disease may be tumour cell invasion and metastasis, breast cancer, colorectal cancer, gastric carcinoma, gynaecological cancer, lung cancer, oesophageal cancer, prostate cancer, scirrhous cancer, tissue regeneration.

The agent may be used for the preparation of a medicament and may be used with a pharmaceutically acceptable carrier.

The present invention will now be described by way of Example only with reference to the accompanying drawings, which are briefly described below:-

Figure 1. Characterization of anti-p120 and 2B12 immunoreactivity. Various cell lines and a primary culture of bovine brain endothelial cells (Brain EC) were lysed in SDS sample buffer and analyzed by SDS-PAGE followed by immunoblotting with anti-p120 antibody (Panel A) or the 2B12 antibody (Panel B). The exposure times were 1 minute for Panel A (apart from that for the MDBK cells which was 10 seconds) and 15 minutes for Panel B. The migration of p120 (o) and p100 (•) has been indicated.

Figure 2. p120 protein is recognized by the anti-p120 antibody and cross-reacts with the 2B12 antibody, whereas p100 is recognized only by anti-p120. MDBK cells were lysed in TDS buffer. Immunoprecipitations were performed using the anti-p120 antibody or 2B12 followed by cross-blotting (Panel A). MDBK cells were chosen because p120 and p100 are well separated by SDS-PAGE and react with both antibodies. In Panel B, MDBK cells were labelled with [<sup>35</sup>S]methionine, lysed in TDS buffer and then immunoprecipitated using anti-p120 or 2B12. Proteins were separated by SDS-PAGE and detected by autoradiography. Clearly, the broad bands (Panel A and see Fig. 1 A) corresponding to p120 and p100 as detected by immunoblotting are resolved as multiple bands. The same bands as seen in the 2B12 immunoprecipitate (o) are seen in the anti-p120 immunoprecipitates. In the anti-p120 immunoprecipitates, additional bands (•) corresponding to p100 are also observed.

Figure 3. Comparison of anti-p120 and anti-E-cadherin immunoprecipitates from MDCK and Caco-2 cells. Panel A: MDCK cells were labelled with [<sup>35</sup>S]methionine and then lysed in either TX buffer or TDS buffer. Immunoprecipitations were performed using anti-p120 or anti-E-cadherin (rr1). Proteins were separated by SDS-PAGE followed by fluorography. Bands corresponding to E-cadherin (E),  $\alpha$ - ( $\alpha$ ) and  $\beta$ -catenin ( $\beta$ ) which are seen strongly in the rr1 immunoprecipitates have been indicated. The major anti-p120 reactive band migrates at approximately the position of that of p100 (see Fig. 1 A). 2B12 does not appear to react with canine protein (see Fig. 1 B) making positive identification of p120 difficult. Panel B: Caco-2 cells were labelled with [<sup>35</sup>S]methionine and then lysed in TX buffer. Immunoprecipitations were performed using anti-E-cadherin (HECD-1), anti- $\beta$ -catenin or anti-p120 antibodies. Proteins were separated by SDS-PAGE followed by fluorography. In all cases, four major comigrating bands were immunoprecipitated, corresponding in order of increasing mobility to E-cadherin,  $\alpha$ -,  $\beta$ - and  $\gamma$ -catenin.

Figure 4. Detection of anti-p120 reactive material in anti-E-cadherin immunoprecipitates and E-cadherin in the anti-p120 immunoprecipitates. MDCK cells were lysed in TX buffer. Immunoprecipitations were performed with either antibody to E-cadherin (rr1), the anti-p120 antibody or anti-focal adhesion kinase (anti-FAK). A sham immunoprecipitation was performed in the absence of primary antibody (-). Following separation by SDS-PAGE, parallel blots were probed using: rr1 (arrowhead, E-cadherin); anti- $\beta$ -catenin (arrowhead,  $\beta$ -catenin); anti-p120 (arrowhead, anti-p120 reactivity); anti-FAK (arrowhead, FAK). Clearly, rr1 and the anti-p120 antibody immunoprecipitated E-cadherin and  $\beta$ -catenin. rr1 could also immunoprecipitate anti-p120 immunoreactive material, but to a lesser extent than that immunoprecipitated by the anti-p120 antibody. Anti-FAK could only immunoprecipitate FAK, and the sham immunoprecipitation did not result in any detectable E-cadherin,  $\beta$ -catenin or anti-p120 reactive material.

Figure 5. Comparison of anti- $\beta$ -catenin, anti-p120 and 2B12 immunoprecipitates from primary cultures of bovine brain endothelial cells. Cells were labelled with [ $^{35}$ S]methionine and then lysed in TX buffer or TDS buffer. Immunoprecipitations were performed using anti- $\beta$ -catenin, anti-p120 and 2B12. Proteins were separated by SDS-PAGE followed by fluorography. In the  $\beta$ -catenin immunoprecipitation from TX lysed cells, bands corresponding to a cadherin (C),  $\alpha$ -catenin ( $\alpha$ ) and  $\beta$ -catenin ( $\beta$ ) have been indicated. Under TDS conditions, a band (o) corresponding to p120 is seen in the anti-p120 and 2B12 immunoprecipitates and a further band (\*) is only seen in the anti-p120 immunoprecipitate, corresponding to p100.

Figure 6. Detection of p120/p100 in  $\beta$ -catenin immunoprecipitates and  $\beta$ -catenin in anti-p120 immunoprecipitates from human umbilical vein endothelial cells (Panel A) and ECV304 cells (Panel B). Cells were lysed in either TX or TDS buffer. Lysates were immunoprecipitated using anti- $\beta$ -catenin or the anti-p120 antibody, followed by analysis by SDS-PAGE and immunoblotting. The migration of  $\beta$ -catenin (arrowheads), p100 (•) and p120 (○) have been indicated. Note the absence of anti-p120 reactive material in the  $\beta$ -catenin immunoprecipitates obtained using TDS buffer. Even though  $\beta$ -catenin and p100, as defined by its reactivity with anti-p120 antibody, migrate very closely, they are clearly distinct proteins.

Figure 7. Localization of anti-p120 reactivity and  $\beta$ -catenin in MDCK cells, and brain endothelial cells. MDCK cells (Panel a,b) and porcine brain endothelial cells (Panel c,d) were co-labelled with anti-p120 antibody (Panel a,c) and anti- $\beta$ -catenin antibody (Panel b,d). Secondary antibodies were fluorescein-conjugated anti-mouse and rhodamine-conjugated anti-rabbit. In this instance, it was verified that each secondary antibody was absolutely specific for its designated species of primary antibody. Bar: 20  $\mu$ m.

Fig. 8. Distribution of anti-p120 immunoreactivity in brain and skeletal muscle tissue of the rat. Brain tissue was co-labelled with the anti-p120 antibody (a,c) and anti- $\alpha$ -catenin (b,d). Anti-p120 immunoreactivity and  $\alpha$ -catenin co-localise at intercellular junctions of choroid plexus epithelium (arrows, a and b) and ventricular ependymal cells (arrowheads, a and b). Both antigens also co-localise at interendothelial junctions of blood vessels of macrovascular origin (arrows in c and d). Microvascular profiles in brain sections were identified by labelling with anti-collagen IV antibody (f); co-labelling with the anti-p120 antibody (e), revealed the presence of antigen at interendothelial junctions (e, arrows). In these microvessels,  $\alpha$ -catenin co-localised with anti-p120 immunoreactivity (not shown). In muscle tissue which had been cut perpendicular to the orientation of the muscle fibres, anti-p120 immunoreactivity is limited to areas between muscle fibres where blood vessels are located (g, arrows). Higher magnification reveals a punctate staining pattern (h, arrows) which is likely to reflect anti-p120 immunoreactivity at interendothelial junctions. In Panels a,c and e, bp depicts the brain parenchyma; in Panel a, v the ventricular lumen and cp the choroid plexus; in Panel g, m refers to muscle tissue. Bars: (a, g), 100  $\mu$ m; (c, e, h), 25  $\mu$ m.

Fig. 9. Sequence analysis of peptides derived by LysC proteolysis of p100 from Caco-2 cells. Caco-2 cells (20 x confluent 9 cm dishes) were lysed in TX buffer to minimize nuclear lysis. Insoluble protein was removed by centrifugation. Deoxycholate and SDS were added to the supernatant to give (w/v) 0.5 % and 0.2 % final concentration, respectively. Addition of these detergents results in dissociation of p100 and p120 from the cadherin/catenin complex. p120 and p100 were immunoprecipitated from the lysate using the anti-p120 antibody (which also recognizes p100) from Transduction Laboratories, rabbit anti-mouse IgG and Protein A Sepharose. The immune complex was washed five times and then dissociated by the addition of Laemmli sample buffer followed by heating at 100°C for 5 minutes. Proteins were precipitated by addition of four volumes of ethanol and incubation at -20°C for 16 hours. The precipitate was resolved by SDS-PAGE (6 % acrylamide) and proteins were visualized by Coomassie Blue. Protein corresponding to p100 was excised from the gel and digested with LysC. Peptides were separated by HPLC and sequenced. Mouse p120 sequence was described by Reynolds et al., 1992. Clearly, human p100 is closely related to mouse p120.

In this figure the standard single letter amino acid code is used. Additionally "X" is used to designate any amino acid.

5        In the Examples given later the following materials and methods were used:-

#### **Antibodies**

10        The anti-canine E-cadherin antibody rr1 developed by Gumbiner and Simons (1986) was provided by Barry Gumbiner (Memorial Sloan-Kettering Cancer Center, NY) or obtained from the Developmental Studies Hybridoma Bank maintained by the Department of Pharmacology and Molecular Sciences, 15 Johns Hopkins University School of Medicine, Baltimore, MD 21205, and the Department of Biological Sciences, University of Iowa, Iowa City, IA 52242, under contract N01-HD-2-3144 from the NICHD. The anti-human E-cadherin antibody HECD-1 (Shimoyama et al., 1989) was from Takara Biomedicals (Shiga, Japan). Anti-p120 and anti-focal adhesion kinase (FAK) antibodies were from Transduction Laboratories (Lexington, KY). The anti-p120 antibody 2B12 (Kanner et al., 1990) was a gift from J.T. Parsons (University of Virginia, Charlottesville, VA). The peptide-directed antibodies against  $\alpha$ - and  $\beta$ - catenin (Staddon et al., J. Cell Sci. in press) were kindly provided by Kurt Herrenknecht (Eisai London Research)

Laboratories Ltd., University College London, London UK). All secondary antibodies used for immunoprecipitation and immunocytochemistry were from Jackson Laboratories Inc. (West Grove, PA). HRP-conjugated secondary antibodies used for immunoblotting were from Amersham (Buckinghamshire, UK).

### Cells

The following cells were cultured at 37°C in medium containing 100 U/ml penicillin and 100 µg/ml streptomycin: Caco-2 (epithelial cells derived from a human colonic tumour: 5 % CO<sub>2</sub>, MEM, 10 % FCS, 1 % non-essential amino-acids, 1 µg/ml insulin); ECV304 (a cell line derived from human umbilical vein endothelial cells: 5 % CO<sub>2</sub>, M199, 10 % FCS); LLC-PK1 (epithelial cells derived from porcine kidney: 5 % CO<sub>2</sub>, M199, 10 % FCS); MDBK (epithelial cells derived from bovine kidney: 5 % CO<sub>2</sub>, MEM, 10 % FCS); Strain I MDCK cells (epithelial cells derived from canine kidney: 5 % CO<sub>2</sub>, MEM, 10 % FCS); RBE4 cells (immortalized rat brain endothelial cells (see Durieu-Trautmann et al., 1993): 5 % CO<sub>2</sub>, α-MEM: Ham's F10 (1:1), 10 % FCS, 0.3 mg/ml geneticin, 1 ng/ml bFGF); Swiss 3T3 fibroblasts (10 % CO<sub>2</sub>, DMEM, 10 % FCS). Caco-2, ECV304, LLC-PK1 and MDBK cells were obtained from the European Collection of Animal Cell Cultures (Salisbury, UK). MDCK cells were provided by Barry Gumbiner. RBE4 cells were from Pierre Couraud (Université Paris VII, Paris, France) and Swiss 3T3 fibroblasts were from Enrique Rozengurt (Imperial Cancer Research Fund, London, UK). Human umbilical vein endothelial cells were from Clonetics (Palo Alto, CA) and cultured according to the manufacturer's instructions. Primary cultures of bovine or porcine brain endothelial cells were grown as described by Rubin et al. (1991). For experimental purposes, confluent cultures of Caco-2, MDBK, MDCK and brain endothelial cells were established on tissue culture-treated, polycarbonate Transwell filters (polycarbonate, 0.4 µm; Costar, Cambridge, MA). Other cells were grown on tissue culture plastic.

### Immunoblotting and Immunoprecipitation

Whole cell lysates from cultures maintained for 16-20 hours in 0.5 % serum were prepared by rapidly replacing the medium with hot Laemmli sample buffer (Laemmli, 1970) supplemented with 5 mM EDTA, followed by heating at 100°C for 5 minutes. Proteins were resolved by slab-gel electrophoresis as described by Laemmli, (1970). The gels were equilibrated in buffer containing 48 mM Tris, 39 mM glycine, 0.03 % SDS (w/v) and 20 % methanol (v/v), and then transferred to nitrocellulose filters (Hybond ECL, Amersham). After Ponceau S staining, the filters were blocked in 5 % (w/v) non-fat dried milk in PBS at 4°C for 16-18 hours. Filters were then incubated with primary antibody in PBS containing 0.05 % Tween-20 and 1 % BSA, followed by detection with appropriate HRP-conjugated secondary antibody and chemiluminescence (ECL, Amersham).

Immunoprecipitations were performed at 4°C. Cultures were rinsed with PBS and then lysed in either TX buffer (1 % (v/v) Triton X-100, 25 mM Hepes, 2 mM EDTA, 0.1 M NaCl, 25 mM NaF, 1 mM vanadate, 25 µM phenylarsine oxide, pH 7.6 (adjusted with NaOH), 1 mM PMSF, 10 µg/ml soybean trypsin inhibitor, 0.1 U/ml α<sub>2</sub>-macroglobulin, 10 µg/ml leupeptin) or TDS buffer, which was identical to the TX buffer except that it was supplemented with 0.5 % (w/v) sodium deoxycholate and 0.2 % (w/v) SDS. The cells were incubated with lysis buffer for 10-15 minutes and then scraped. The lysates were collected and centrifuged at 14,000 x g for 20 minutes. The supernatant was precleared with Protein A Sepharose (Pharmacia, UK) for 1-2 hours and then incubated with primary antibody for 1 hour followed by a further 1 hour with Protein A Sepharose alone, in the case of rabbit antibodies, or together with rabbit anti-mouse antibodies for the mouse monoclonal antibodies. After five washes in lysis buffer, immune complexes were dissociated by addition of Laemmli sample buffer followed by heating at 100°C for 5 minutes. Protein analysis was by SDS-PAGE and immunoblotting as described above.

For [<sup>35</sup>S]methionine labelling, the cultures were washed twice in methionine-free MEM supplemented with 0.5 % FCS. The cells were incubated for 16-18 hours in this medium containing 50  $\mu$ Ci/ml [<sup>35</sup>S]methionine (>1000 Ci/mmol, Amersham). Protein analysis was by SDS-PAGE, followed by fixation in 15 % methanol/10 % acetic acid. Labelled protein was detected either by direct autoradiography at room temperature or by fluorography at -80°C following impregnation of the gel with Amplify (Amersham).

#### Immunocytochemistry

Cells were fixed at room temperature for 15 minutes in 3 % paraform-aldehyde made up in PBS containing 0.5 mM CaCl<sub>2</sub> and 0.5 mM MgSO<sub>4</sub>. Fixed cells were washed and then permeabilized by incubation with 0.5 % Triton X-100 in PBS for 10 minutes. After washing, the cells were incubated for 30 minutes in PBS containing 10 % calf serum and 0.1 M lysine, pH 7.4. Incubation with primary antibody was in PBS containing 10 % calf serum for 1 hour. After washing, the cells were then incubated for 30-60 minutes with a 1:100 dilution of fluorophore-conjugated anti-mouse or anti-rabbit IgG, as appropriate, in PBS containing 10 % calf serum. After washing, the filters were mounted with Citifluor (Citifluor Products, Canterbury, UK) and examined using a Nikon Microphot-FXA fluorescence microscope fitted with 40 x and 60 x objectives. Photographs were taken using Kodak T-MAX film (400 ASA).

For the preparation of cryosections, brain and skeletal muscle from CO<sub>2</sub>-asphyxiated rats were removed and rapidly frozen in liquid nitrogen. Tissue blocks were mounted in Tissue Tek (R. Lamb, London, UK) and sections of 5-10  $\mu$ m thickness were cut on a Bright cryostat, air-dried and stored for up to 4 weeks at -20°C. After thawing, the sections were fixed and permeabilized as described above. The sections were then washed, blocked with PBS containing 10% calf serum for 15 minutes and incubated with primary antibody diluted in

PBS containing 10% calf serum for 2 hours. After washing, they were incubated with PBS containing 10% calf serum with either 10% goat serum or 10% donkey serum, as appropriate for the host of the secondary antibody, for 15 minutes. They were then incubated with secondary antibody diluted in PBS and serum for 1 hour. Sections were washed, mounted and examined as described above.

#### EXAMPLE 1

##### Antibody characterization: p120 and p100 proteins

SDS lysates of a variety of cell types were analyzed by immunoblotting for expression of p120 using an anti-p120 antibody (Fig. 1 A). This antibody was raised against a fusion protein containing amino acids 790-911 at the C-terminus of p120. The antibody clearly reacted with two distinct bands of apparent molecular mass 120 kDa and 100 kDa in extracts of Caco-2, ECV 304, RBE4, Swiss 3T3, LLC-PK1, MDCK and brain endothelial cells. We will hereafter refer to these two bands as p120 and p100. With the MDCK cells a broad continuous band was detected. The relative intensity of the staining of p120 and p100 depended on the cell type. For example, p100 was more intense than p120 in Caco-2 cells and in the Swiss 3T3 cells the reverse was observed. These blots were also probed with the monoclonal anti-p120 antibody CB12, raised against a phosphotyrosine immunoprecipitate prepared from pp60<sup>src</sup> transformed chicken embryo fibroblasts (Kanner et al., 1990). This antibody recognised only the upper of the two bands (Fig. 1 B). CB12 reacted strongly with bovine protein, moderately with rodent protein and weakly with human protein. However, it failed to react with extracts of the LLC-PK1 (porcine) and MDCK (canine) cells, reflecting limited cross-species reactivity.

The relationship between the known pp60<sup>src</sup> substrate p120 and p100 was further examined by immunoblot analysis of immunoprecipitates from cells lysed

in TDS buffer, used to minimize protein-protein interaction (Fig. 2 A). MDBK cells were used since both the anti-p120 and 2B12 antibodies worked well (Fig. 1). In the anti-p120 immunoprecipitates, p120 and p100 were detected by blotting with the anti-p120 antibody. 2B12 only reacted with p120 in these immunoprecipitates. In 2B12 immunoprecipitates, blotting with both 2B12 and the anti-p120 antibody revealed p120. Thus, 2B12 recognized p120, whereas the anti-p120 antibody recognized the same p120 protein as 2B12 and, in addition, a p100 protein.

Autoradiographic analysis of immunoprecipitates of [<sup>35</sup>S]methionine-labelled MDBK cells, as resolved by SDS-PAGE (Fig. 2 B), revealed that the p120 protein migrated as a cluster of bands that were identical in the both 2B12 immunoprecipitates and the anti-p120 immunoprecipitates. However, in the anti-p120 immunoprecipitates a similar cluster of additional bands corresponding to p100 was detected. Technically, detection by immunoblotting was light based (chemiluminescence) which does not have the greater resolution of direct autoradiography, hence the multiple bands seen in Fig. 2 B appear as a broad band in Fig. 2A. The basis of the multiplicity of the bands corresponding to p120 and p100 in the MDBK cells is not clear. However, [<sup>32</sup>P]phosphate-labeling of these cells raised the possibility that these bands may represent differentially phosphorylated protein (results not shown).

#### Association with the cadherin/catenin complex: epithelial cells

We next examined the possibility of association of p120 or p100 with other proteins. When TDS lysates of [<sup>35</sup>S]methionine-labelled MDBK cells were immunoprecipitated with the anti-p120 antibody, p120 and p100 were isolated as the major proteins corresponded to those detected by immunoblotting (Fig. 2). Similarly, the major bands seen in immunoprecipitates from TDS lysates of [<sup>35</sup>S]methionine-labelled MDCK cells corresponds to the broad band detected by immunoblotting (cf. Fig. 1 A and Fig. 3 A). In contrast, the anti-p120

immunoprecipitates from MDCK cells lysed in TX buffer, used to preserve macromolecular protein complexes, revealed additional distinct bands at approximately 130 kDa, 105 kDa and 97 kDa (Fig. 3 A). Proteins of similar molecular mass were seen in E-cadherin immunoprecipitates from similarly lysed cells (Fig. 3 A). In this case, the 130 kDa band corresponds to E-cadherin, the 97 kDa band to  $\beta$ -catenin, and the 105 kDa band to  $\alpha$ -catenin, which is dissociated from the complex by lysis in TDS buffer (see McCrea and Gumbiner, 1991). Thus, it appeared that the anti-p120 antibody may be capable of immunoprecipitating the cadherin/catenin complex.

Caco-2 cells were also [ $^{35}$ S]methionine-labelled, lysed in TX buffer and immunoprecipitations were performed using the anti-E-cadherin antibody HECD-1, anti- $\beta$ -catenin antibody or the anti-p120 antibody. HECD-1 or anti- $\beta$ -catenin clearly immunoprecipitated four major bands corresponding in order of increasing mobility to E-cadherin,  $\alpha$ -,  $\beta$ -, and  $\gamma$ -catenin. Similar bands were immunoprecipitated, but to a lesser extent, with the anti-p120 antibody (Fig. 3 B). These results (not shown) were also obtained in MDBK cells when  $\beta$ -catenin immunoprecipitates were compared with those obtained with the anti-p120 antibody. Thus, it appears that the anti-p120 antibody can immunoprecipitate proteins that comigrate with those of the cadherin/catenin complex from a variety of epithelial cell lines.

The comigration of bands in the anti-p120 and anti-E-cadherin immunoprecipitates suggests, but does not prove, that anti-p120 is capable of immunoprecipitating the cadherin/catenin complex. Therefore, the identity of the proteins in the E-cadherin and anti-p120 immunoprecipitates was examined by immunoblotting (Fig. 4). As expected, E-cadherin and  $\beta$ -catenin were both present in the rr1 immunoprecipitates. An anti-p120 reactive band displaying a mobility that was similar to that of  $\beta$ -catenin was also present in the rr1 immunoprecipitates. In the anti-p120 immunoprecipitates, anti-p120 reactive material was

also present, unsurprisingly, but to a greater extent and migrated as a broader band than that seen in the rrl immunoprecipitates. In the anti-p120 immunoprecipitates, E-cadherin and  $\beta$ -catenin also were clearly detected. Omission of primary antibody did not result in E-cadherin, anti-p120 reactivity or  $\beta$ -catenin in the immunoprecipitates. Immunoprecipitation with anti-FAK antibody, of the same species and isotype as the anti-p120 antibody, immunoprecipitated FAK but not E-cadherin,  $\beta$ -catenin or anti-p120 reactivity (Fig. 4).

It should be noted that although  $\beta$ -catenin and the anti-p120 immunoreactive band seen in the anti-E-cadherin immunoprecipitates have a similar mobility (Fig. 4), this does not represent cross-reactivity of the anti-p120 antibody with  $\beta$ -catenin. Thus, in  $\beta$ -catenin immunoprecipitates from MDCK cells lysed in TDS buffer, which resulted in dissociation of anti-p120 reactive material from the complex of  $\beta$ -catenin and E-cadherin (see Fig. 3 A), the anti-p120 antibody failed to react with  $\beta$ -catenin. Conversely, anti-p120 immunoprecipitates from TDS lysed cells fail to contain  $\beta$ -catenin immunoreactivity (results not shown). We also observed the anti-p120 immunoreactive band, as seen in the rrl immunoprecipitates (Fig. 4), in  $\alpha$ -catenin and  $\beta$ -catenin immunoprecipitates (results not shown).

Therefore, on the basis of the comigration of [ $^{35}$ S]methionine-labelled bands seen in anti-E-cadherin and anti-p120 immunoprecipitates of cells lysed in TX buffer, it appeared that the anti-p120 antibody could immunoprecipitate the cadherin/catenin complex. By immunoblotting, the unequivocal identification of E-cadherin and  $\beta$ -catenin in the anti-p120 immunoprecipitates verified this interpretation.

### Association with the cadherin/catenin complex; endothelial cells

Endothelial cells form functional complexes of similar composition to those found in epithelial cells (for review, see Rubin, 1992). As shown (Fig. 1), endothelial cells also contain p120 and p100 proteins. We therefore asked whether p100/p120 is also associated with the cadherin/catenin complex in endothelial cells. Brain endothelial cells were labelled with [<sup>35</sup>S]methionine, lysed with TX buffer and immunoprecipitated with either anti- $\beta$ -catenin antibody, anti-p120 antibody or, because of its strong reactivity against bovine protein (see Fig. 1), 2B12. The anti- $\beta$ -catenin antibody clearly immunoprecipitated three major proteins (Fig. 5), corresponding in order of increasing mobility to a 120-130 kDa band, probably representing a cadherin, and  $\alpha$ - (105 kDa) and  $\beta$ -catenin (97 kDa). The anti-p120 antibody immunoprecipitated proteins (but to a lesser extent than those observed in the  $\beta$ -catenin immunoprecipitates) that comigrated with  $\alpha$ - and  $\beta$ -catenin and a broad 120 kDa band, representing p120 (see later), that migrated between the presumptive cadherin and  $\alpha$ -catenin (Fig. 5). In the 2B12 immunoprecipitates, proteins that comigrated with  $\alpha$ - and  $\beta$ -catenin were also observed but to a much lesser extent than observed in the  $\beta$ -catenin immunoprecipitates. Again, a broad 120 kDa band was observed in these immunoprecipitates. From TDS lysates, anti- $\beta$ -catenin antibody could still precipitate  $\beta$ -catenin, but anti-p120 antibody failed to do so. Instead, a diffuse band was observed that migrated slightly more slowly than  $\beta$ -catenin, and this was absent in the 2B12 immunoprecipitates, therefore representing p100. In the 2B12 immunoprecipitates, only the 120 kDa band was observed. Thus, association between the catenins and p120 or p100 appeared to be observed in TX lysates, but not TDS lysates, of brain endothelial cells.

To explore the generality of these observations, similar experiments were performed using human umbilical vein endothelial cells (Fig. 6 A) and ECV304 cells (Fig. 6 B) but using immunoblot analysis. Thus, from cells lysed in TX buffer,

anti- $\beta$ -catenin antibody immunoprecipitated  $\beta$ -catenin, as expected, but also two bands of anti-p120 immunoreactivity, corresponding to p100 and p120. Similarly, the anti-p120 antibody immunoprecipitated p100 and p120, as expected, and  $\beta$ -catenin. However, as for the brain endothelial cells and the epithelial cells, TDS buffer resulted in dissociation of  $\beta$ -catenin and p100/p120. These data corroborate those obtained from the analysis of the immunoprecipitates from [ $^{35}$ S]-methionine-labelled brain endothelial cells (Fig. 5).

### EXAMPLE 3

#### Localization to cell-cell junctions

To investigate the cellular localization of p100/p120, immunocytochemical analysis was performed. Unfortunately, CB12 did not label cells under the fixation conditions that were tried (paraformaldehyde and methanol). However, the anti-p120 antibody (which recognizes p120 and p100) stained cells well. Double-labelling of MDCK cells with anti- $\beta$ -catenin and the anti-p120 antibody revealed a striking co-localization of  $\beta$ -catenin and anti-p120 immunoreactivity (Fig. 7 a,b). Similar results were found for the brain endothelial cells (Fig. 7 c,d).

Immunocytochemistry was further performed on frozen sections of brain and skeletal muscle (Fig. 8). Anti-p120 immunoreactivity co-localized with  $\alpha$ -catenin at intercellular junctions of choroid plexus epithelium and ventricular ependymal cells (Fig. 8 a,b). In addition, the anti-p120 antibody stained interendothelial junctions in both large (Fig. 8 c,d) and small blood vessels of the brain (Fig. 8 e,f) and blood vessels of muscle tissue (Fig. 8 g,h). At the given detection level, anti-p120 immunoreactivity was limited to endothelial and epithelial cells in the two types of tissue investigated.

In the present study, immunoblot analysis of a variety of different cells revealed that the anti-p120 monoclonal antibody 2B12 (Kanner et al., 1990) recognized a broad band of approximately 120 kDa. Another anti-p120 monoclonal antibody that had been raised against the C-terminal portion of p120 recognized the same bands as 2B12 and, in addition, another cluster of bands at 100 kDa. The reason for the multiplicity of bands at 120 kDa and 100 kDa is not clear, but obviously they are immunologically related. The pattern of appearance of these multiple bands also depended on the cell type. For example in MDCK cells, the bands were very diffuse, whereas in MDBK cells they were clearly resolved, especially when visualized by [<sup>35</sup>S]methionine labelling in the absence of fluorographic reagent. It is possible that post-translational modification, such as phosphorylation and myristylation may account for the appearance of multiple bands. The relationship, apart from the immunological one, between p120 and p100 is not yet clear. The Northern blots described by Reynolds et al. (1992) suggested the possibility of p120-related gene products or alternatively spliced transcripts. Furthermore, Southern analysis apparently indicated that one or more p120-related genes exist (Reynolds et al., 1992). It follows that the cluster of bands corresponding to p120 and the similar cluster to p100 could also represent splice variants of, respectively, p100 and p120. It is also possible that p100 could simply represent a degradation product of p120, although samples were prepared in denaturing buffer and immunoprecipitations were performed in the presence of inhibitors of a broad spectrum of proteases.

Our results provide evidence that p120/p100 associates with the cadherin/catenin complex. Thus, from TX-solubilized epithelial cells, anti-p120 antibody immunoprecipitated [<sup>35</sup>S]methionine-labelled proteins that comigrated with those immunoprecipitated by cadherin antibodies (Fig. 3). Immunoblot analysis verified the identity of these proteins as cadherins and catenins (Fig. 4).

Similarly, anti-p120 reactive protein was seen in the anti-E-cadherin immunoprecipitates from MDCK cells, although on the basis of mobility this appears to represent primarily p100, not p120 (Fig. 4). However, it should be noted that p120 immunoreactive protein was not unequivocally identified in the MDCK cells because of the poor reactivity of the 2B12 antibody with canine protein. This apparent association between p120/p100 and the cadherin/catenin complex was not restricted to epithelial cells as similar results were obtained using endothelial cells, both by [<sup>35</sup>S]methionine-labelling (Fig. 5) and immunoblot analysis (Fig. 6). However, although our results clearly show an interaction between catenins and p120/p100 in endothelial cells, we did not identify a cadherin in the anti-p120 immunoprecipitates. The interpretation of the biochemical analyses of the protein complexes in the epithelial and endothelial cells is further supported by the immunocytochemical study. In cultured cells and in tissue sections, anti-p120 immunoreactivity was strongly associated with intercellular junctions, displaying a localization that was remarkably similar to that obtained with anti- $\beta$ -catenin antibody (Figs. 7 and 8). Here, we were restricted to using the anti-p120 antibody which might recognize either p120 or p100, or both.

Of course, one explanation for the ability of anti-p120 antibody to immunoprecipitate the cadherin/catenin complex is that it cross-reacts with one of the known components of the complex. However, this is unlikely for the following reasons. TDS lysis of cells dissociates  $\alpha$ -catenin and  $\gamma$ -catenin from the cadherin and  $\beta$ -catenin complex (see McCrea and Gumbiner, 1991). In anti-E-cadherin immunoprecipitates from MDCK cells lysed in TDS buffer, anti-p120 antibody failed to react with E-cadherin or the co-precipitating  $\beta$ -catenin. In anti- $\alpha$ -catenin immunoprecipitates from MDCK cells lysed in TDS buffer, anti-p120 antibody failed to react with  $\alpha$ -catenin (results not shown). As shown in Fig. 6,  $\beta$ -catenin immunoprecipitates from TDS-lysed endothelial cells, although clearly

containing  $\beta$ -catenin, and presumably a cadherin, did not contain any anti-p120 reactivity. Cross-reactivity with  $\gamma$ -catenin, a protein of approximately 85 kDa, is unlikely as the immunoblots shown in Fig. 1 fail to show reactivity with protein below 100 kDa.

One problem that has to be addressed concerns the lack of any obvious [ $^{35}$ S]methionine-labelled bands that correspond to p120/p100 in the cadherin immunoprecipitates from TX lysed cells. Thus, as shown in Fig. 3, rr1 and the anti-p120 immunoprecipitates from MDCK cells look remarkably similar. However, it is clear from the anti-p120 immunoprecipitates from TDS lysed cells that the anti-p120 reactive material migrates as a broad band, the bulk of which migrates between  $\alpha$ - and  $\beta$ -catenin, with the remainder migrating slightly slower than  $\alpha$ -catenin. In conjunction with the immunoblot analysis shown in Fig. 4, which demonstrates that only a fraction of the anti-p120 immunoreactive material is immunoprecipitated by rr1, and the fact that p120 contains about half of the number of methionines in  $\alpha$ - and  $\beta$ -catenin, if we assume steady-state labelling, then it is obvious that it would be difficult to see any labelling corresponding to the anti-p120 immunoreactive material in the rr1 immunoprecipitates. Similarly, with respect to the experiments with the brain endothelial cells (Fig. 5), under the extraction conditions employed it appears that only a fraction of the pool of the cadherin/catenin complex associates with p100, and even less with p120. Thus, it would be difficult to see labelling of bands corresponding to p120 and p100 in the  $\beta$ -catenin immunoprecipitates as this region of the gel is already occupied by other major bands. It is also difficult to discern a band corresponding to the cadherin seen in the  $\beta$ -catenin immunoprecipitates in either the anti-p120 or 2B12 immunoprecipitates because of the dominance of the labelling of p120 (Fig. 5).

We emphasize that there are also clear quantitative differences with respect to the ability of 2B12 and anti-p120 antibody to immunoprecipitate the cadherin/catenin complex. Due to poor reactivity of 2B12 with canine protein, this

could not be addressed with MDCK cells. In bovine brain endothelial cells, the anti-p120 antibody immunoprecipitated the cadherin/catenin complex, although not as well as anti- $\beta$ -catenin antibody. 2B12 was even less effective than anti-p120. As anti-p120 recognizes p120 and p100, and 2B12 only p120, the difference in the amount of the complex immunoprecipitated must be attributable to greater association of p100 with the catenins. As far as we can tell, the efficiency of anti-p120 and 2B12 in immunoprecipitation was similar in this experiment (see Fig. 5 for the similar intensity of a band corresponding to p120 in the anti-p120 and 2B12 immunoprecipitates from cells lysed in TX buffer). These data also indicate that independent complexes of p100 and p120 exist with catenins, rather than a catenin/p100/p120 complex. If the latter were the case, 2B12 would be expected to immunoprecipitate as much of the catenins as anti-p120 antibody.

Our study provides a link between p120/p100 and adherens junction proteins. It is possible that such an interaction, perhaps via the influence of regulatory kinases, such as src, lyn and yes (see Tsukita et al., 1991), may play a role in the modulation of cadherin function, and thereby other cellular functions influenced by the adherens junction. With respect to phosphorylation, the tyrosine phosphatase inhibitor phenylarsine oxide was found to cause an increase in tight junction permeability in MDCK cells (Staddon et al., *J. Cell Sci.* in press). This inhibitor increased the tyrosine phosphorylation of the anti-p120 immunoreactive material (a major p100 band, a minor p120 band) in these cells, as analyzed by anti-p120 immunoblotting of anti-p120 immunoprecipitates from SDS lysates (results not shown; see Staddon et al., *J. Cell Sci.* in press). The p120/p100 proteins could also be involved in the interaction between cadherins and the actin-based cytoskeleton. p120/p100 may also be part of a signalling cascade, communicating information about the state of cell-cell adhesiveness to the interior of the cell.

$\beta$ -catenin is an arm protein (McCrea et al., 1991) and can associate with cadherins and the APC gene product (Rubinfeld et al., 1993; Su et al., 1993), also

an arm protein (see Peifer et al., 1994). p120 is an arm protein (Reynolds et al., 1992; Peifer et al., 1994), and, as we describe here, p100 is an immunologically related protein. These proteins can interact with  $\beta$ -catenin. The exact nature of the interaction between p120/p100 and the catenins remains to be established. These proteins may interact directly, or associate with different regions of the cytoplasmic domain of cadherins. Other linking or intermediary binding proteins could also be involved. Clearly, there appears to be diverse interactions among arm proteins, suggesting the importance of the arm motif in intracellular signalling.

In summary, our studies identify p100 as a p120-related protein. We present evidence that these proteins interact with the cadherin/catenin complex. Given the important role of the cadherin/catenin complex in cellular transformation and the identification of p120 as a pp60<sup>src</sup> substrate, this suggests that p120/p100 may play a role in cellular growth control and other processes, such as tight junction permeability control, via an influence on cell-cell adhesion.

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CLAIMS

1. A protein:

5 (a) having a molecular weight of about 100 kDa and being associated in vivo with the catenin/cadherin complex of epithelial or endothelial cells, which is referred to herein as p100, or

(b) having substantial amino acid homology with p100, or

10 (c) being a fragment of p100 or of a protein having substantial amino acid homology with p100.

2. A protein according to claim 1 which comprises one or more of the amino acid sequences given below:

15 (i) NISFGRDQDNK

(ii) HAIPNLV

(iii) XVLINK

(iv) XPIEDPANDTVDFPX

(v) XPSGALRNLAVDARX,

20 wherein the letters used are based on the standard single letter amino acid code, but wherein "X" designates any amino acid.

25 3. A nucleic acid molecule coding for a protein according to claim 1 or claim 2, or being complementary to a nucleic acid molecule according to claim 1 or claim 2.

30 4. A vector comprising a nucleic acid molecule according to claim 3.

5. A host cell comprising a vector according to claim 4.

The use of a protein according to claim 1 or claim of p120 for studying cell-cell adhesion.

The use of a protein according to claim 1 or claim of p120 for studying tight junction permeability.

An agent which alters or blocks association between or p120 and the catenin/cadherin complex of epithelial or endothelial cells for use in the treatment diagnosis of a disease.

An agent according to claim 8, wherein the disease selected from: tumour cell invasion and metastasis, breast cancer, colorectal cancer, gastric carcinoma, gynaecological cancer, lung cancer, oesophageal cancer, prostate cancer, scirrhous cancer, tissue regeneration or developmental disorders.

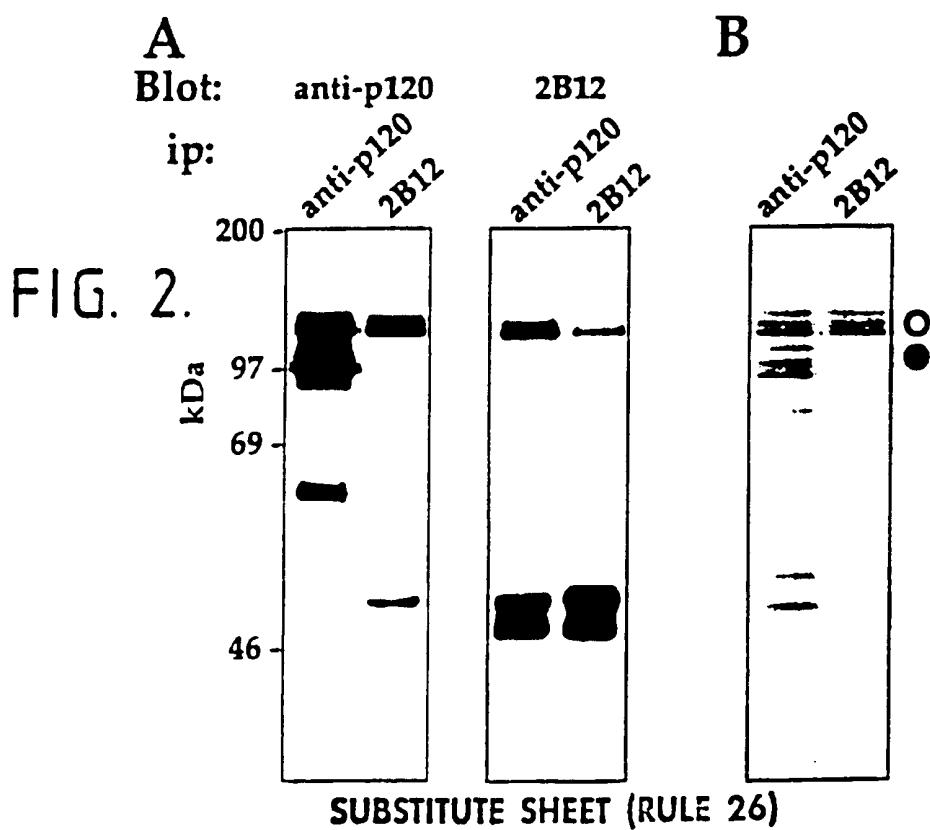
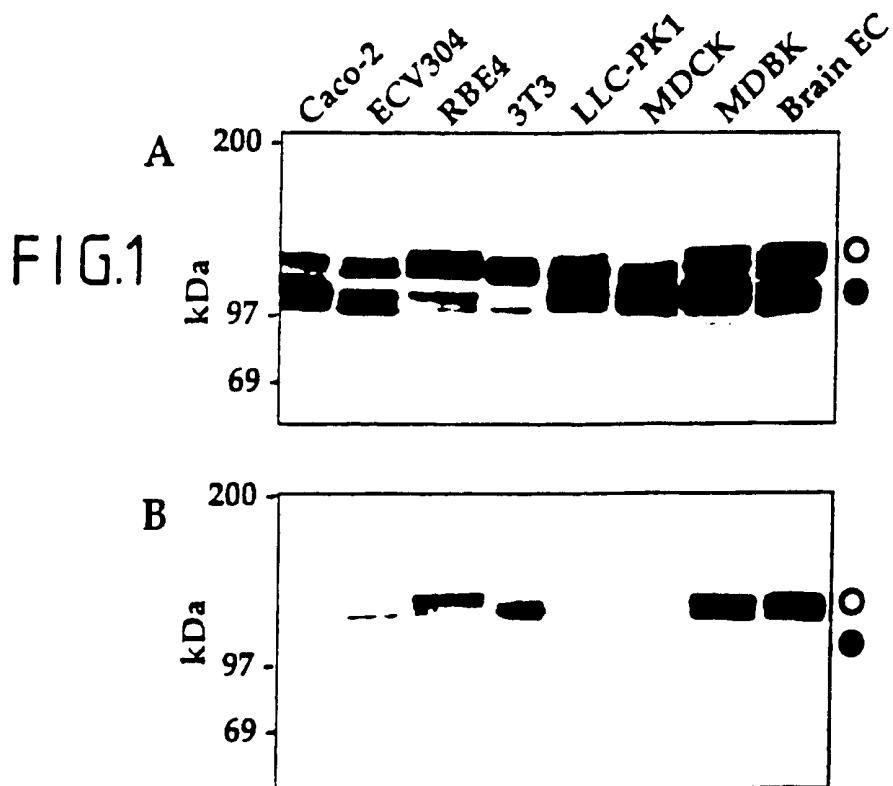
An agent according to claim 8 or claim 9 which is a peptide.

The use of an agent according to any of claims 8 to 2 the manufacture of a medicament for the treatment of disease selected from: tumour cell invasion and metastasis, breast cancer, colorectal cancer, gastric carcinoma, gynaecological cancer, lung cancer, oesophageal cancer, prostate cancer, scirrhous cancer, tissue regeneration or developmental disorders.

30

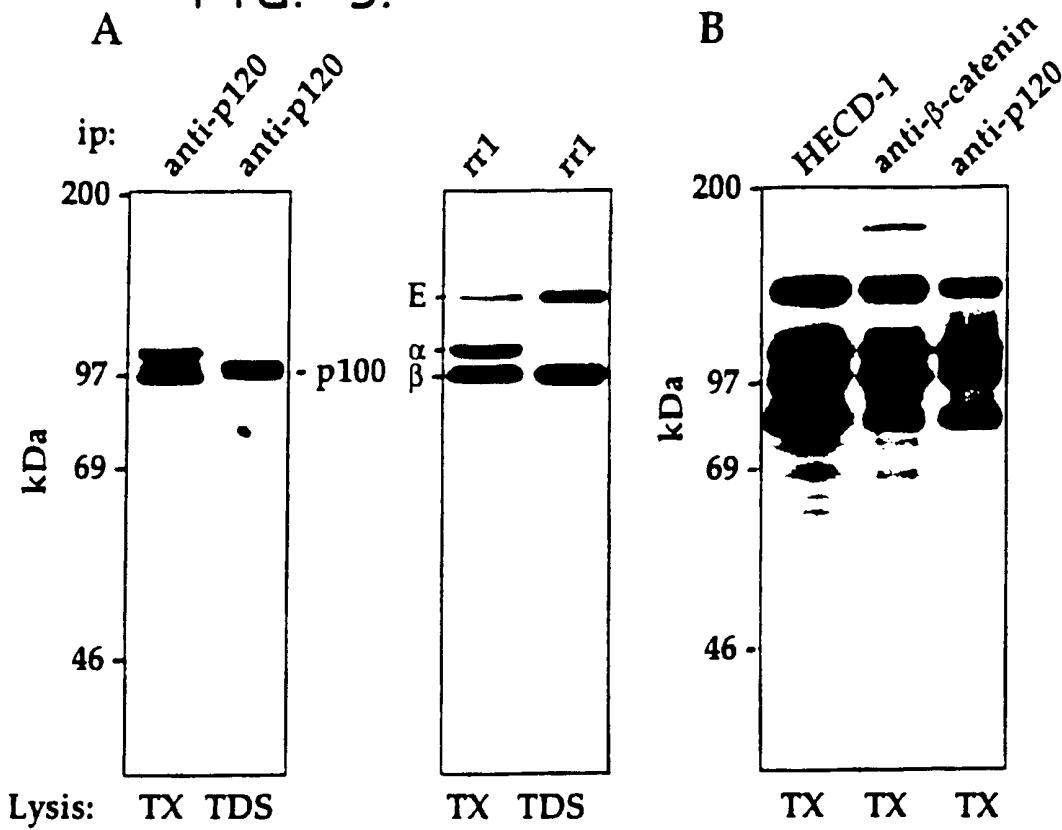
The protein p100, substantially as hereinbefore described, with reference to the accompanying examples.

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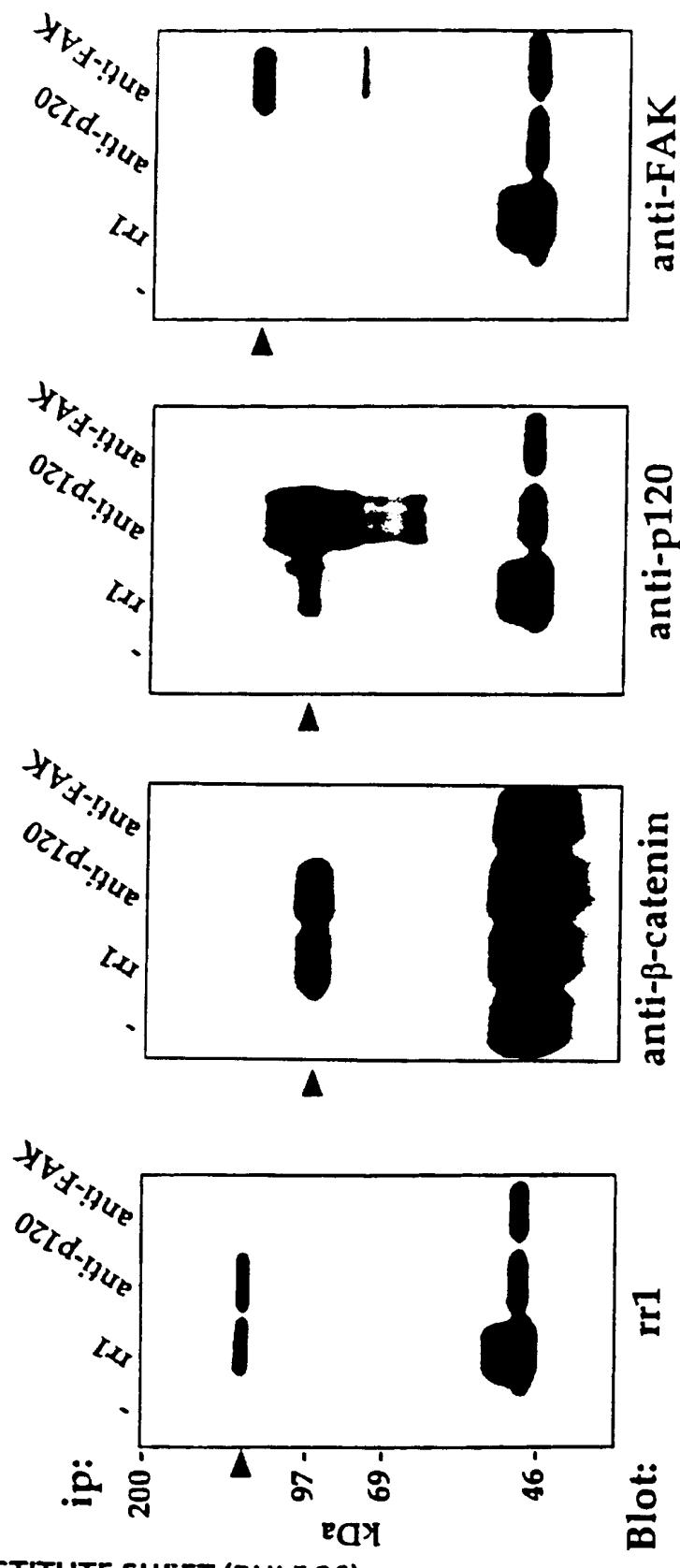
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FIG. 3.



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FIG. 4.



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FIG. 5. ip:

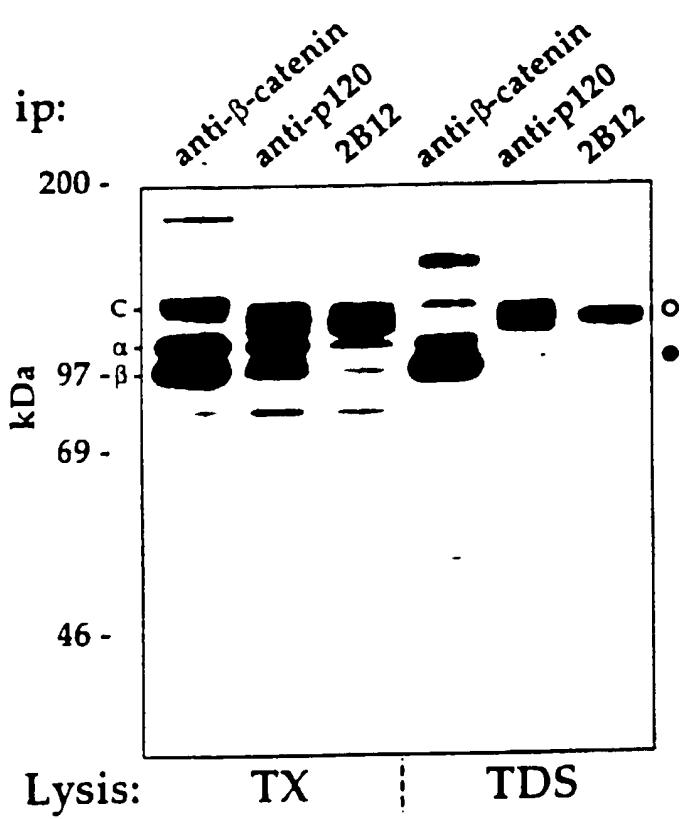
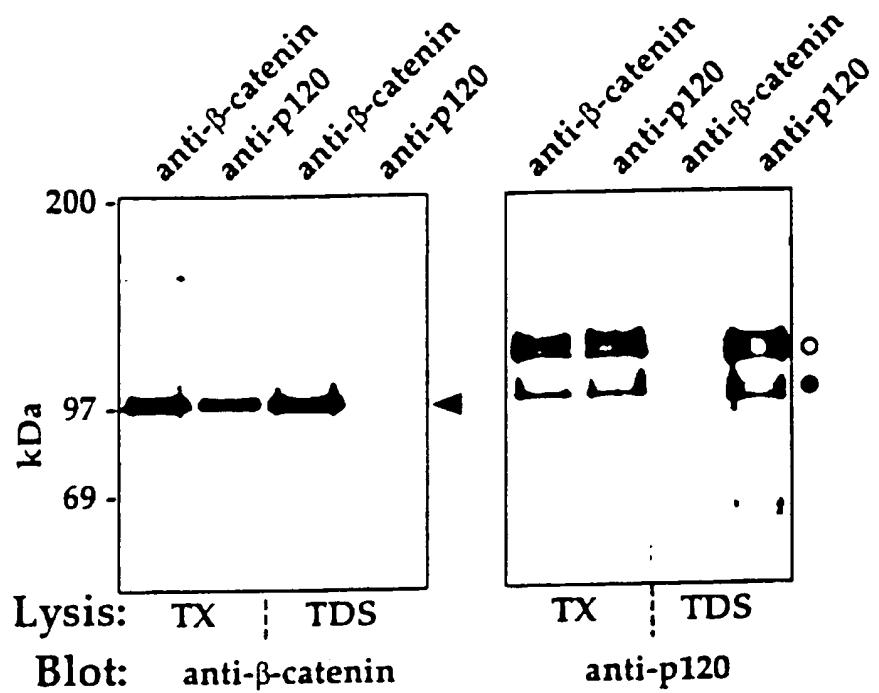
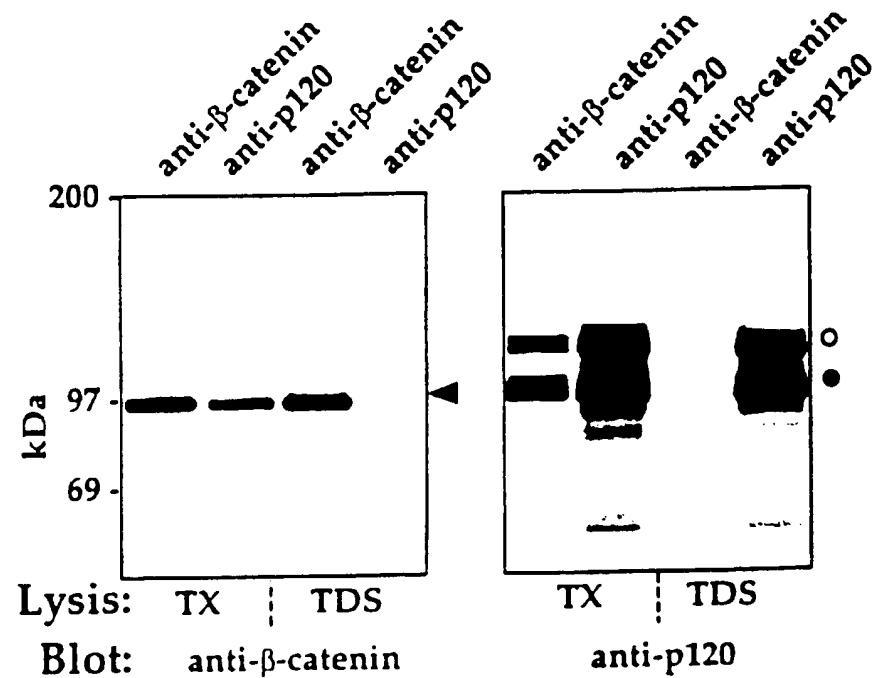
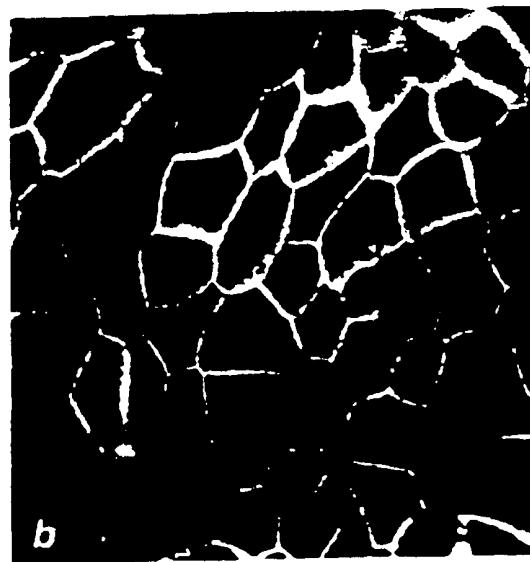
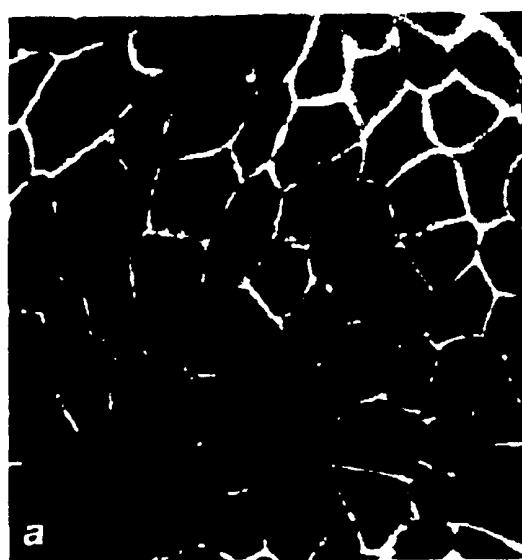


FIG. 6.

**A****B**

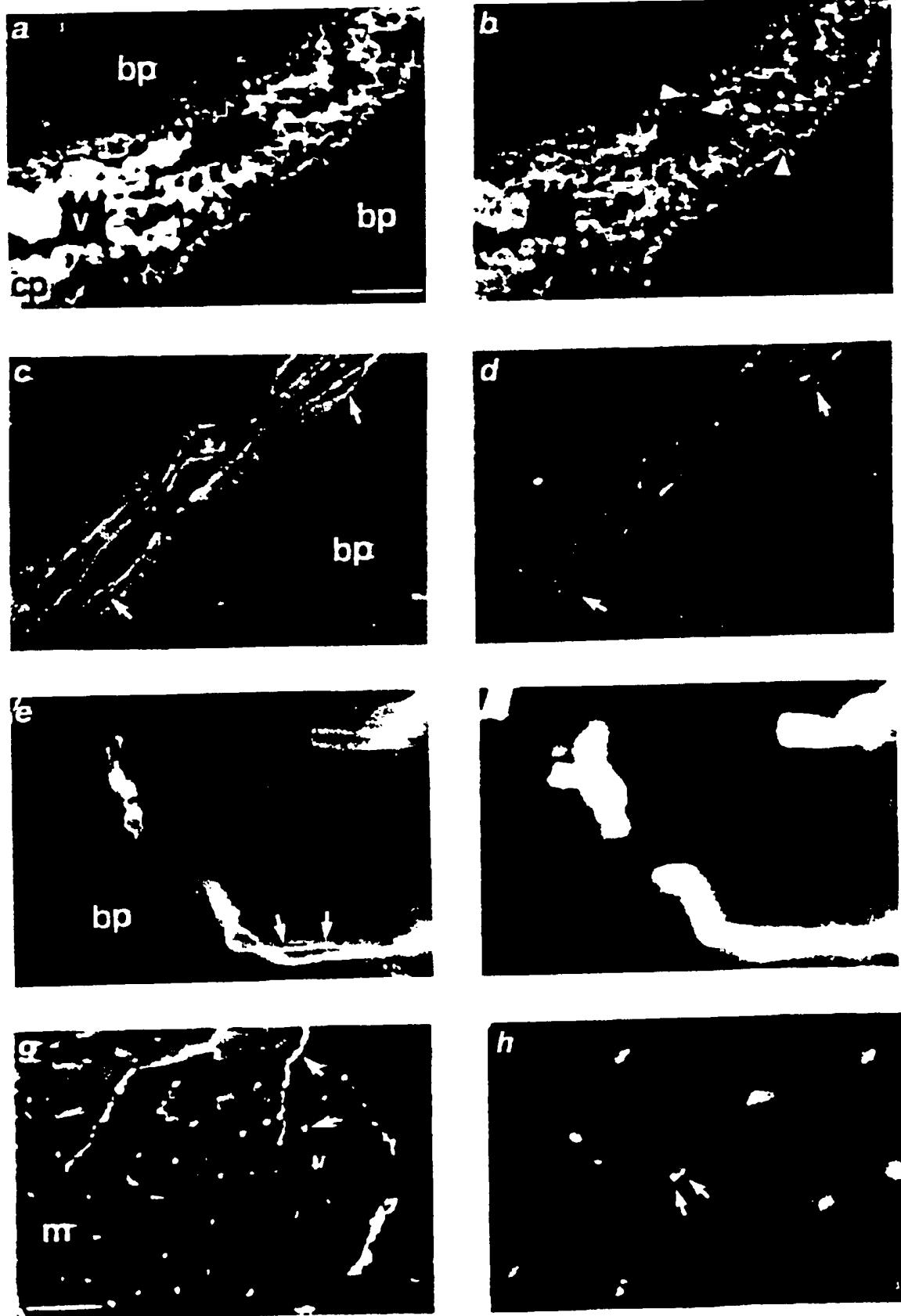
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FIG. 7.



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FIG. 8.



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FIG. 9.

SEQUENCER RESULTS	PUBLISHED SEQUENCE	RESIDUES
NISFGRDQDNK	NISFGRDQDNK	434-444
HAIPNLV	HARPNLV	744-751
XVLINK	LVLINK	799-804
XPIEDPANDTVDFPX	KPTEDPANDTVDFPX	629-642
XPSGALRNLAVDARX (Tentative)	AASGALRNLAVDARK	723-738

PCT

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International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification <sup>6</sup> : <b>C12N 15/12, 15/85, 5/10, C07K 14/47, 16/18, C12Q 1/00, A61K 39/395</b></p>		A3	<p>(11) International Publication Number: <b>WO 96/16170</b> (43) International Publication Date: <b>30 May 1996 (30.05.96)</b></p>
<p>(21) International Application Number: <b>PCT/GB95/02686</b> (22) International Filing Date: <b>16 November 1995 (16.11.95)</b></p>		<p>(81) Designated States: JP, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).</p>	
<p>(30) Priority Data: <b>9423372.3 18 November 1994 (18.11.94) GB</b></p>		<p><b>Published</b> <i>With international search report.</i></p>	
<p>(71) Applicant (<i>for all designated States except US</i>): EISAI CO., LTD. [JP/JP]; 4-6-10, Koishikawa, Bunkyo-ku, Tokyo 112- 88 (JP).</p>		<p>(88) Date of publication of the international search report: <b>1 August 1996 (01.08.96)</b></p>	
<p>(72) Inventor; and (75) Inventor/Applicant (<i>for US only</i>): STADDON, James, Michael [GB/GB]; 44 Oakwood Park Road, Southgate, London N14 6QA (GB).</p>			
<p>(74) Agents: WHITE, Martin, P. et al.; Kilburn &amp; Strode, 30 John Street, London WC1N 2DD (GB).</p>			
<p>(54) Title: PROTEINS INVOLVED IN THE REGULATION OF CELL-CELL ADHESION AND USES THEREOF</p>			
<p>(57) Abstract</p>			
<p>A p100 protein, which has a molecular weight of about 100 kDa and appears to be related to p120, is involved in the regulation of cell-cell adhesion.</p>			

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GA	Gabon				

# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/GB 95/02686

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC 6	C12N15/12	C12N15/85	C12N5/10	C07K14/47	C07K16/18
	C12Q1/00	A61K39/395			

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)  
IPC 6 C07K C12N C12Q A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>ONCOGENE (1992), 7(12), 2439-45 CODEN: ONCNES; ISSN: 0950-9232, 1992, XP002002165</p> <p>REYNOLDS, ALBERT B. ET AL: "P120, a novel substrate of protein tyrosine kinase receptors and of p60v-src, is related to cadherin -binding factors. beta.- catenin, plakoglobin and armadillo" cited in the application see abstract see page 2439, right-hand column, paragraph 3 - page 2441, left-hand column, paragraph 3 see page 2444, left-hand column, paragraph 2</p> <p>---</p> <p>-/-</p>	1,3-7,12

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

\* Special categories of cited documents :

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- 'T' later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
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- 'A' document member of the same patent family

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Date of the actual completion of the international search	Date of mailing of the international search report
6 May 1996	10.05.96
Name and mailing address of the ISA  European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+ 31-70) 340-2040, Tx. 31 651 epo nl, Fax (+ 31-70) 340-3016	Authorized officer  Montero Lopez, B

## INTERNATIONAL SEARCH REPORT

International Application No  
PCT/GB 95/02686

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>CELL, vol. 65, no. 5, May 1991, NA US, pages 849-857, XP002002166 AKIRA NAGAFUCHI ET AL.: "The 102 kd Cadherin-associated protein: Similarity to vinculin and posttranscriptional regulation of expression" cited in the application see the summary see page 849, right-hand column, last paragraph - page 850, left-hand column, paragraph 1 see page 851, left-hand column, paragraph 2 - paragraph 3; figure 3 see page 853, left-hand column, paragraph 2</p> <p>---</p>	1,3-7,12
A	<p>J. CELL SCI. (1994), 107(8), 2259-70 CODEN: JNCSAI; ISSN: 0021-9533, 1994, XP002002167 HATZFELD, MECHTHILD ET AL: "Band 6 protein, a major constituent of desmosomes from stratified epithelia, is a novel member of the armadillo multigene family" see the summary see page 2262, left-hand column, paragraph 2 - page 2267, left-hand column, paragraph 2</p> <p>---</p>	1-7,12
A	<p>PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF USA, vol. 87, no. 9, May 1990, WASHINGTON US, pages 3328-3332, XP002002168 STEVEN B. KANNER ET AL.: "Monoclonal antibodies to individual tyrosine-phosphorylated protein substrates of oncogene-encoded tyrosine kinases" cited in the application see abstract see page 3328, right-hand column, paragraph 2</p> <p>---</p> <p>-/-</p>	8

## INTERNATIONAL SEARCH REPORT

International Application No  
PCT/GB 95/02686

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	<p>J. CELL BIOL. (1995), 130(2), 368-81      CODEN: JCLBA3; ISSN: 0021-9525,      1995, XP002002169</p> <p>STADDON, JAMES M. ET AL: "p120, a p120-related protein ( p100 ), and the cadherin / catenin complex"      see abstract      see page 370, left-hand column, paragraph 3      see page 371, right-hand column, last paragraph - page 372, left-hand column, paragraph 1      see page 371, right-hand column, paragraph 2 - page 374, right-hand column, paragraph 2      see page 379, left-hand column, last paragraph - page 380, right-hand column, paragraph 2      ---</p> <p>MOLECULAR AND CELLULAR BIOLOGY,      vol. 14, no. 12, December 1994, WASHINGTON      US,      pages 8333-8342, XP000561742</p> <p>ALBERT B. REYNOLDS ET AL.:      "Identification of a new catenin: the tyrosine kinase substrate p120cas associates with E-cadherin complexes"      see abstract      see page 8334, right-hand column, last paragraph - page 8336, left-hand column, paragraph 1      see page 8341, left-hand column, paragraph 3      -----</p>	1,2,6,7, 12
P,X		6

# INTERNATIONAL SEARCH REPORT

International application No.

PCT/GB95/02686

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
  
2.  Claims Nos.: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
  
3.  Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

- claims 1-5, 12 and, partially, 6-11
- claims 6-11 partially

- see next sheet -

1.  As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
  
2.  As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
  
3.  As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
  
4.  No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

### Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/SA/210

- Protein P100, nucleic acid encoding it, vector and host cell containing the same, uses thereof and agent blocking its association to the catenin/cadherin complex
- Use of protein P120 for studying cell-cell adhesion and agent blocking the association of P120 with the catenin/cadherin complex